Kazuhiro R Nitta

List of Publications by Year
in descending order

Source: https://exaly.com/author-pdf/8689498/publications.pdf

Version: 2024-02-01

22	3,932	566801 15	713013 21
papers	citations	h-index	g-index

23	23	23	7420
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Neonatal-onset mitochondrial disease: clinical features, molecular diagnosis and prognosis. Archives of Disease in Childhood: Fetal and Neonatal Edition, 2022, 107, 329-334.	1.4	9
2	Development of Leigh syndrome with a high probability of cardiac manifestations in infantile-onset patients with m.14453GÂ›ÂA. Mitochondrion, 2022, 63, 1-8.	1.6	2
3	Fatal Perinatal Mitochondrial Cardiac Failure Caused by Recurrent De Novo Duplications in the ATAD3 Locus. Med, 2021, 2, 49-73.e10.	2.2	33
4	Clinical heterogeneity in patients with m.4412Gâ€¯›â€¯A MT-TM mutation and different heteroplasmy levels. Mitochondrion, 2021, 59, 214-215.	1.6	0
5	Genome sequencing and RNAâ€seq analyses of mitochondrial complex I deficiency revealed Alu insertionâ€mediated deletion in NDUFV2. Human Mutation, 2021, 42, 1422-1428.	1.1	4
6	Long-term prognosis and genetic background of cardiomyopathy in 223 pediatric mitochondrial disease patients. International Journal of Cardiology, 2021, 341, 48-55.	0.8	14
7	Diverse Mechanisms of Resistance to Decitabine and Venetoclax Therapy in Newly Diagnosed and Relapsed/Refractory AML Inferred By Transcriptome Analysis. Blood, 2021, 138, 2244-2244.	0.6	2
8	ANISEED 2019: 4D exploration of genetic data for an extended range of tunicates. Nucleic Acids Research, 2020, 48, D668-D675.	6.5	30
9	Functional annotation of human long noncoding RNAs via molecular phenotyping. Genome Research, 2020, 30, 1060-1072.	2.4	109
10	A homozygous variant in <scp>NDUFA8</scp> is associated with developmental delay, microcephaly, and epilepsy due to mitochondrial complex I deficiency. Clinical Genetics, 2020, 98, 155-165.	1.0	18
11	High-Throughput Protein Production Combined with High- Throughput SELEX Identifies an Extensive Atlas of Ciona robusta Transcription Factor DNA-Binding Specificities. Methods in Molecular Biology, 2019, 2025, 487-517.	0.4	15
12	ANISEED 2017: extending the integrated ascidian database to the exploration and evolutionary comparison of genome-scale datasets. Nucleic Acids Research, 2018, 46, D718-D725.	6.5	90
13	The interaction landscape between transcription factors and the nucleosome. Nature, 2018, 562, 76-81.	13.7	259
14	Impact of cytosine methylation on DNA binding specificities of human transcription factors. Science, 2017, 356, .	6.0	912
15	Myt1l safeguards neuronal identity by actively repressing many non-neuronal fates. Nature, 2017, 544, 245-249.	13.7	180
16	DNA-dependent formation of transcription factor pairs alters their binding specificity. Nature, 2016, 534, S15-S16.	13.7	280
17	DNA-dependent formation of transcription factor pairs alters their binding specificity. Nature, 2015, 527, 384-388.	13.7	462
18	Conservation of transcription factor binding specificities across 600 million years of bilateria evolution. ELife, 2015, 4, .	2.8	316

#	Article	IF	Citations
19	<scp>H</scp>em<scp>R</scp> is an <scp>O</scp>mp<scp>R</scp>/<scp>P</scp>ho<scp>B</scp>â€like response regulator from <scp>L</scp>eptospira, which simultaneously effects transcriptional activation and repression of key haem metabolism genes. Molecular Microbiology, 2014, 94, 340-352.	1.2	23
20	DNA-Binding Specificities of Human Transcription Factors. Cell, 2013, 152, 327-339.	13.5	1,085
21	Expression of Sox1 during Xenopus early embryogenesis. Biochemical and Biophysical Research Communications, 2006, 351, 287-293.	1.0	37
22	XSIP1 is essential for early neural gene expression and neural differentiation by suppression of BMP signaling. Developmental Biology, 2004, 275, 258-267.	0.9	48